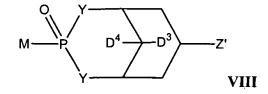
## IN THE CLAIMS:

Please cancel claims 58-149, 161, 163-164, and 167-170 without prejudice. The Applicants reserve the right to pursue claims to non-elected inventions in a Divisional application.

Please amend the claims to read as follows:

- (Once amended) The compounds of claim 1 wherein V is -H, and Z is selected from the group consisting of -CHR<sup>2</sup>OH, -CHR<sup>2</sup>OC(O)R<sup>3</sup>, and -CHR<sup>2</sup>OCO<sub>2</sub>R<sup>3</sup>.
- 35. (Once amended) The compounds of claim 22 wherein W and W' are independently selected from the group consisting of -H, alkyl, aralkyl, alicyclic, aryl, substituted aryl, heteroaryl, and substituted heteroaryl.
  - 173. (Once amended) The compounds of claim 1 that are of formula VIII:



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wherein:

Z' is selected from the group consisting of -OH,  $-OCO_2R^3$ ,  $-OC(O)R^3$ , and  $-OC(O)SR^3$ ;

R<sup>2</sup> is selected from the group consisting of R<sup>3</sup> and -H;

R<sup>3</sup> is selected from the group consisting of alkyl, aryl, alicyclic, and aralkyl;

R<sup>6</sup> is selected from the group consisting of -H, and lower alkyl, acyloxyalkyl, alkoxycarbonyloxy alkyl and lower acyl;

one Y is -O- and the other Y is -NR<sup>6</sup>-;

 $D^3$  is -H;

D<sup>4</sup> is selected from the group consisting of -H, alkyl, -OH, -OR<sup>2</sup> and -OC(O)R<sup>3</sup>.

M is selected from the group that attached to PO<sub>3</sub><sup>2-</sup>, P<sub>2</sub>O<sub>6</sub><sup>3-</sup>, P<sub>3</sub>O<sub>9</sub><sup>4-</sup> or P(O)(NHR<sup>6</sup>)O is a biologically active agent but is not an FBPase inhibitor, and is attached to the phosphorus in formula I via a carbon, oxygen, sulfur or nitrogen atom;

with the provisos that:

1) M is not –NH(lower alkyl), -N(lower alkyl)<sub>2</sub>, -NH(lower alkylhalide),

-N(lower alkylhalide)<sub>2</sub>, or -N(lower alkyl) (lower alkylhalide); and SAN/62125.3.

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2) R<sup>6</sup> is not lower alkylhalide; and pharmaceutically acceptable prodrugs and salts thereof.